

This Listing of Claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS

1. *(currently amended)*: A method for producing a population of mononuclear cells overexpressing IL-10 polypeptide which cells (i) are not selected on the basis of specificity for a predetermined antigen and (ii) treat an inflammatory disease or condition in an antigen-independent manner, the method comprising:
 - (a) ~~providing mammalian peripheral blood mononuclear cells wherein, if the cells comprise lymphocytes, the lymphocytes are not selected or enriched on the basis of their antigen specificity;~~
 - (a[[b]]) modifying introducing into at least a portion fraction of a mammalian peripheral blood said mononuclear cell[[s]] population,
among which cells lymphocytes are not selected or enriched on the basis of antigen specificity,
by introducing into said cells an expression construct that comprises a nucleotide sequence encoding an IL-10 polypeptide; and,
 - (b[[c]]) recovering, from said modified mononuclear cells, cells that
 - (i) overexpress the IL-10 polypeptide, and
 - (ii) are not specific for a predetermined antigen,thereby producing said population of mononuclear cells overexpressing IL-10 that are capable of treating said inflammatory disease.
2. *(currently amended)*: A method according to claim 1, wherein the expression construct is introduced into an enriched fraction or subset of said peripheral blood mononuclear cell[[s]] population are provided.
3. *(currently amended)*: A method according to claim 2, wherein the enriched fraction or subset is selected from the group consisting of (i) lymphocytes ~~or a subset thereof~~, (ii) macrophages, (iii) monocytes [or] and (iv) dendritic cells (DC).
4. *(currently amended)*: A method according to claim 1, wherein prior to the introducing step (a[[b]]), the mononuclear cells are induced to, or allowed to, proliferate.
5. *(previously presented)*: A method according to claim 4, wherein the mononuclear cells are induced to proliferate by a proliferating agent.

6. *(previously presented)*: A method according to claim 5, wherein the proliferating agent is one or more of

- (a) an anti-CD3 antibody;
- (b) an anti-CD28 antibody; or
- (c) phytohemagglutinin.

7. *(currently amended)*: A method according to claim 1, wherein subsequent to step (a[[b]]), the modified mononuclear cells are fractionated to yield an enriched cell [[a]] fraction or subset thereof.

8. *(previously presented)*: A method according to claim 7, wherein the fraction or subset comprises enriched

- (i) lymphocytes or a subset thereof,
- (ii) macrophages or monocytes, or
- (iii) dendritic cells.

9. *(currently amended)*: A method according to claim 1, wherein subsequent to step (a[[b]]), the modified mononuclear cells are enriched for cells that overexpress the IL-10-encoding nucleotide sequence.

10. *(withdrawn; previously presented)*: A method for producing a pharmaceutical composition comprising mononuclear cells overexpressing IL-10, which method comprises

- (a) producing the mononuclear cells overexpressing IL-10 in accordance with claim 1, and
- (b) combining said cells with an acceptable pharmaceutical carrier.

11. *(withdrawn; currently amended)*:: A composition comprising mononuclear cells that are not selected to be specific for a predetermined antigen which cells are modified to ~~that~~ comprise an IL-10 transgene.

12. *(withdrawn; currently amended)*: A T lymphocyte composition that is a fraction of the mononuclear cells according to claim 11, ~~wherein the mononuclear cells comprise which~~ T cells ~~that~~ comprise said IL-10 transgene.

13. *(withdrawn; currently amended)*: A T lymphocyte composition according to claim 12, wherein the T cells functionally mimic regulatory T cells in that they inhibit:

- (a) proliferation of autologous responder cells, and/or
- (b) production of pro-inflammatory cytokine IL-12 by dendritic cells.

14. *(withdrawn)*: A pharmaceutical composition comprising the composition according to claim 11, and a pharmaceutically acceptable carrier.
15. *(withdrawn)*: A method of treating a disease or condition associated with undesired activation and/or expansion of T cells, which method comprises administering an effective amount of a pharmaceutical composition according to claim 14 to a subject suffering from said disease or condition.
16. *(withdrawn)*: A method according to claim 15, wherein the disease or condition is a Th1-mediated disease or condition.
17. *(withdrawn)*: A method according to claim 16, wherein the Th1-mediated disease or condition is Crohn's disease, reactive arthritis, insulin-dependent diabetes, colitis, pancreatitis, an inflammatory lung disease, an inflammatory eye disease, multiple sclerosis, Hashimoto's thyroiditis, Graves' disease, chronic articular rheumatism, contact dermatitis, psoriasis, graft rejection, graft-versus-host disease, or sarcoidosis.

Claims 18 and 19 [previously canceled]

20. *(currently amended)* A method according to claim 3 wherein the enriched lymphocyte subset comprises an enriched population of B lymphocytes eells, T lymphocytes eells or CD4+ lymphocytes eells.
21. *(currently amended)* A method according to claim 8 wherein the enriched lymphocyte[[s]] subset comprises an enriched population of B lymphocytes eells, T lymphocytes eells or CD4+ lymphocytes eells.

CANCEL Claim 22

23. *(withdrawn)* A method according to claim 16 wherein the Th1-mediated disease is an inflammatory disease.